

=> fil hcap

FILE 'HCAPLUS' ENTERED AT 18:09:22 ON 11 MAY 2007

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

FILE COVERS 1907 - 11 May 2007 VOL ISS ISS

FILE LAST UPDATED: 10 May 2007 (20070510/ED)

held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications.

The CA Lexicon is the copyrighted intellectual property of the the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 11 May 2007 VOL 146 ISS 21

FILE LAST UPDATED: 1 May 2007 (20070501/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate

=> d his nofile

(FILE 'HOME' ENTERED AT 17:10:24 ON 11 MAY 2007)

FILE 'HCAPLUS' ENTERED AT 17:10:40 ON 11 MAY 2007

L1 1 SEA ABB=ON PLU=ON US2004214088/PN

FILE 'REGISTRY' ENTERED AT 17:13:19 ON 11 MAY 2007

L2 43 SEA ABB=ON PLU=ON (105-58-8/BI OR 107-31-3/BI OR

FILE 'HCAPLUS' ENTERED AT 17:30:06 ON 11 MAY 2007

L3 QUE ABB=ON PLU=ON PORO? OR MICROPORO?

L4 QUE ABB=ON PLU=ON PERMEA? OR PERFORAT? OR SIEVE? OR  
PERVIOUS?

L5 QUE ABB=ON PLU=ON POLYM? OR COPOLYM? OR RESIN? OR  
HOMOPOLYM?

L6 35290 SEA ABB=ON PLU=ON (L3 OR L4) (3A) L5

L7 QUE ABB=ON PLU=ON FILM? OR THINFILM? OR LAYER? OR  
OVERLAY?

L8 5311 SEA ABB=ON PLU=ON L6 (3A) L7

L9 QUE ABB=ON PLU=ON ?MICROSCAL? OR ?MICROSIZ? OR  
?MICROMATERIAL? OR ?MICROCHEM? OR ?MICRO(A) (SCAL? OR  
CHEM? OR SIZ? OR MATERIAL?)

L10 21 SEA ABB=ON PLU=ON L8 AND L9

L11 QUE ABB=ON PLU=ON POLYETHYLENE# OR PE OR POLYPROPYLENE#  
OR PP OR POLYIMIDE# OR PI OR POLYSULFONE# OR POLYSULPHON  
E# OR PSU

L12 QUE ABB=ON PLU=ON POLYURETHANE# OR PUR OR POLYVINYLCHELO  
RIDE# OR POLY(W)VINYLCHELORIDE# OR PVC OR CELLULOSE# OR  
NYLON# OR POLYACRYLONITRILE# OR PAN

L13 QUE ABB=ON PLU=ON POLYVINYLDENE (2A) FLUORIDE# OR  
POLY(W)VINYLDENE (W) FLUORIDE# OR PVDF OR POLYTETRAFLUOROE  
THYLENE# OR PTFE

L14 QUE ABB=ON PLU=ON FIRST? OR 1ST OR 1(W)ST OR PRIMARY?

L15 QUE ABB=ON PLU=ON SECOND? OR 2ND OR 2(W)ND

L16 QUE ABB=ON PLU=ON ELECTROLY?  
 L17 QUE ABB=ON PLU=ON BATTERY  
 L18 20748 SEA ABB=ON PLU=ON (L11 OR L12 OR L13) (L) L16  
 L19 2 SEA ABB=ON PLU=ON L10 AND L16  
 L20 1 SEA ABB=ON PLU=ON L19 AND (L11 OR L12 OR L13)  
 L21 229 SEA ABB=ON PLU=ON L8 AND L18  
 L22 33 SEA ABB=ON PLU=ON L21 AND L14  
 L23 18 SEA ABB=ON PLU=ON L22 AND L15  
 L24 2 SEA ABB=ON PLU=ON L19 OR L20  
 L25 17 SEA ABB=ON PLU=ON L23 NOT L24  
 L26 15 SEA ABB=ON PLU=ON L22 NOT (L24 OR L25)

=> d l24 ibib abs hitstr hitind 1-2

L24 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2004:905471 HCAPLUS Full-text  
 DOCUMENT NUMBER: 141:382157  
 TITLE: Method of fabrication of composite polymer  
           **electrolyte** of different morphologies  
           for lithium secondary battery  
 INVENTOR(S): Lee, Young Gi; Kim, Kwang Man; Ryu, Kwang Sun;  
               Chang, Soon Ho  
 PATENT ASSIGNEE(S): S. Korea  
 SOURCE: U.S. Pat. Appl. Publ., 10 pp.  
           CODEN: USXXCO  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
US 2004214088	A1	20041028	US 2003-748363	200312 29
KR 2004092188	A	20041103	KR 2003-26419	200304 25
JP 2004327422	A	20041118	JP 2003-431458	200312 25
CN 1610169	A	20050427	CN 2003-10125472	200312 31
PRIORITY APPLN. INFO.:			KR 2003-26419	A 200304 25

AB A composite polymer **electrolyte** for a lithium secondary battery and a method of manufacturing the same are provided. The composite polymer **electrolyte** includes a composite film structure which includes a first **porous polymer film** with good mech. properties and a second **porous polymer film** with **submicro-scale** morphol. of more compact porous structure than the first porous polymer structure, coated on a surface of the first **porous polymer film**, and an **electrolyte** solution impregnated into the composite film structure. The different morphologies of the composite film structure enable to an increase in mech. properties and ionic conductivity. Furthermore, the charge/discharge cycle performance and stability of a lithium metal polymer secondary battery are enhanced.

IC ICM H01M010-40  
 INCL 429309000; 429316000; 429317000; 429314000  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 38  
 ST polymer **electrolyte** different morphol lithium secondary  
 battery  
 IT Secondary batteries  
 (lithium; method of fabrication of composite polymer  
**electrolyte** of different morphologies for lithium  
 secondary battery)  
 IT Battery **electrolytes**  
 Composites  
 Polymer morphology  
 (method of fabrication of composite polymer **electrolyte**  
 of different morphologies for lithium secondary battery)  
 IT Acrylic polymers, uses  
 Fluoropolymers, uses  
 Polyamide fibers, uses  
**Polyimides**, uses  
 Polyoxyalkylenes, uses  
**Polysulfones**, uses  
**Polyurethanes**, uses  
 Zeolites (synthetic), uses  
 RL: DEV (Device component use); USES (Uses)  
 (method of fabrication of composite polymer **electrolyte**  
 of different morphologies for lithium secondary battery)  
 IT 96-47-9, 2-Methyltetrahydrofuran 96-48-0,  $\gamma$ -Butyrolactone  
 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate  
 107-31-3, Methyl formate 108-32-7, Propylene carbonate 109-94-4,  
 Ethyl formate 109-99-9, Thf, uses 110-71-4 616-38-6, Dimethyl  
 carbonate 623-53-0, Ethyl methyl carbonate 1344-28-1, Alumina,  
 uses 7631-86-9, Silica, uses 7791-03-9, Lithium perchlorate  
 9002-84-0, **Ptfe** 9002-86-2, Polyvinyl chloride  
 9002-88-4, **Polyethylene** 9003-07-0, **Polypropylene**  
 9003-20-7, Polyvinyl acetate 9003-21-8, Polymethyl acrylate  
 9003-32-1, Polyethyl acrylate 9003-42-3, Polyethyl methacrylate  
 9003-49-0, Polybutylacrylate 9003-63-8, Polybutylmethacrylate  
 9004-34-6, **Cellulose**, uses 9011-14-7, Pmma 9011-17-0,  
 Hexafluoropropylene-vinylidene fluoride copolymer 12003-67-7,  
 Aluminum lithium oxide allio2 13463-67-7, Titania, uses  
 14283-07-9, Lithium tetrafluoroborate 14807-96-6, Talc, uses  
 21324-40-3, Lithium hexafluorophosphate 24937-79-9, **Pvdf**  
 25014-41-9, **Polyacrylonitrile** 25322-68-3, Peo  
 25322-69-4, **Polypropylene** oxide 28960-88-5,  
 Trifluoroethylene-vinylidene fluoride copolymer 33454-82-9,  
 Lithium triflate 90076-65-6  
 RL: DEV (Device component use); USES (Uses)  
 (method of fabrication of composite polymer **electrolyte**  
 of different morphologies for lithium secondary battery)  
 IT 67-64-1, Acetone, uses 67-68-5, DmsO, uses 68-12-2, Dmf, uses  
 872-50-4, n-Methylpyrrolidone, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (method of fabrication of composite polymer **electrolyte**  
 of different morphologies for lithium secondary battery)

L24 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 1978:580757 HCAPLUS Full-text  
 DOCUMENT NUMBER: 89:180757  
 TITLE: Localization of micropores in polymer films  
 AUTHOR(S): Geymayer, W. F.; Grasenick, F.

CORPORATE SOURCE: Zent. Elektronenmikrosk. Graz, Graz, Austria  
 SOURCE: Electron Microsc., Pap. Int. Congr., 9th (1978),  
 Volume 1, 490-1. Editor(s): Sturgess, J. M.  
 Microsc. Soc. Canada: Toronto, Ont.  
 CODEN: 39ERA7  
 DOCUMENT TYPE: Conference  
 LANGUAGE: English

AB The pore channels in a swollen polymer matrix are marked and stabilized for electron microscopic examination by filling the active pore-volume with insol. metal salts or dense metallic ppts. AgCl was precipitated in the micropores of an osmotic membrane by placing a NaCl solution on 1 side and AgNO3 on the other and repeatedly changing the direction of flow. Polymer films coated on metal substrates are investigated by inserting the substrate as a cathode in a suitable **electrolyte** and filling the reactive pore-volume by **electrolytic** separation. After filling the pore channels, the metal deposition widens in a mushroom shape, clearly marking the pore entry.

CC 36-6 (Plastics Manufacture and Processing)

IT **Polymers**, preparation  
 RL: PREP (Preparation)  
 (microporous films, method of marking pores  
 for, for electron microscopy)

IT Pore  
 (micro-, size and location of, in polymer  
 films, method for marking of)

=> d 125 ibib abs hitstr hitind 1-17

L25 ANSWER 1 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2006:1073250 HCAPLUS Full-text  
 DOCUMENT NUMBER: 145:422611  
 TITLE: Composite proton exchange membrane for gas  
 permeation-resistant fuel cell, and its  
 manufacture  
 INVENTOR(S): Mu, Shichun; Chen, Lei; Pan, Mu; Yuan, Runzhang  
 PATENT ASSIGNEE(S): Wuhan University of Technology, Peop. Rep. China  
 SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu,  
 13pp.  
 CODEN: CNXXEV  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Chinese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	---	---	-----	
CN 1707837	A	20051214	CN 2005-10018751	200505 20
PRIORITY APPLN. INFO.:			CN 2005-10018751	200505 20

AB The title membrane is a 3-layered porous polymer reinforced composite exchange membrane having two 1st water-retainable porous polymer reinforced composite exchange membranes as outer layers obtained by compositing inorg. nanoparticles and a 1st solid polymer **electrolyte**, and a 2nd porous polymer reinforced composite exchange membrane as an intermediate layer possessing humidifying and gas resistant functions and obtained by compositing Pt nanoparticles and a 2nd

solid polymer **electrolyte**. The title membrane is manufactured by (1) immersing a porous polymer membrane in ethanol for cleaning and pre-swelling; (2) drying the membrane; (3) preparing 1st solns. of 1st inorg. nanoparticles (SiO<sub>2</sub>, TiO<sub>2</sub>, ZrO<sub>2</sub>, or Zr(HPO<sub>4</sub>)) and 1st solid polymer **electrolytes**; (4) immersing the polymer membrane in the solution of the inorg. nanoparticles and the solid polymer **electrolyte** for 5-20 min; (5) taking the membrane out of the solution; (6) drying the membrane on a heating; (7) roll pressing the membrane; (8) repeating steps (4)-(7) several times to obtain the 1st membrane with required thickness; (9) dispersing Pt nanoparticles in a 2nd solution of 2nd inorg. nanoparticles and a 2nd solid polyelectrolyte; (10) immersing the ethanol treated polymer membrane in the 2nd solution for 5-20 min; (11) repeating steps (4)-(7) several times to obtain the 2nd membrane; (12) sandwiching the 2nd membrane between two 1st membranes, then sandwiching between two PTFE thin films; (13) and hot pressing at 120-135° under 0.5-4 MPa for 1-5 min, and stripping the PTFE thin films.

IC ICM H01M008-02

ICS H01M008-10; H01M002-14; C08J005-22

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

L25 ANSWER 2 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:983911 HCAPLUS Full-text

DOCUMENT NUMBER: 143:289420

TITLE: **Secondary** battery using porous film  
type solvent-free polymer electrolyte filled  
with oligomer/prepolymer electrolyte

INVENTOR(S): Kwak, Seung-Yeop; Jeon, Jae-Deok

PATENT ASSIGNEE(S): Seoul National University Industry Foundation,  
S. Korea

SOURCE: PCT Int. Appl., 26 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005081646	A2	20050909	WO 2005-KR525	20050226
WO 2005081646	A3	20060202		
W:				
AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA,				
CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,				
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP,				
KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX,				
MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE,				
SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,				
VC, VN, YU, ZA, ZM, ZW				
RW:				
BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW,				
AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ,				
DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC,				
NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA,				
GN, GQ, GW, ML, MR, NE, SN, TD, TG				
KR 2005087263	A	20050831	KR 2004-12983	20040226

PRIORITY APPLN. INFO.:

KR 2004-12983

A

20040226

- AB Provided are a solvent-free polymer **electrolyte** and a **secondary** battery employing the same. The solvent-free **polymer electrolyte** includes: a **porous film** including a **first** polymer and a **second** oligomer, the **first** polymer being at least one selected from the group consisting of **poly(vinylidene fluoride-co-hexafluoropropylene)** copolymers, polyvinylidene fluorides, polymethylmethacrylates, **polyacrylonitriles**, polyethyleneoxides, and **celluloses** having a polyether chain and the **second** oligomer being at least one selected from the group consisting of poly(ethylene oxide-co-ethylene carbonate) copolymers with at least one terminal groups substituted by a halogen atom and polyethyleneglycols with at least one terminal groups substituted by a halogen atom; and an **electrolyte** present in the pores of the porous film and including the **second** oligomer and a lithium salt. Since the solvent-free polymer **electrolyte** contains no liquid organic **electrolyte**, it is not accompanied by problems caused by leakage or evaporation of an organic solvent, unlike a gel-type polymer **electrolyte**. Furthermore, the solvent-free polymer **electrolyte** has enhanced ionic conductivity as compared to a conventional solvent-free polymer **electrolyte**.
- IC ICM H01M
- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 38
- IT Polyoxyalkylenes, uses  
RL: DEV (Device component use); USES (Uses)  
(halogenated; **secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)
- IT Fillers  
(inorg.; **secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)
- IT Battery anodes  
Battery electrolytes  
Polymer electrolytes  
**Secondary** batteries  
(**secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)
- IT Carbonaceous materials (technological products)  
Fluoropolymers, uses  
Oligomers  
RL: DEV (Device component use); USES (Uses)  
(**secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)
- IT Zeolites (synthetic), uses  
RL: MOA (Modifier or additive use); USES (Uses)  
(**secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)
- IT 7439-93-2D, Lithium, intercalation compound 7791-03-9, Lithium perchlorate 9004-34-6, **Cellulose**, uses 9011-14-7, Pmma 9011-17-0, Hexafluoropropylene-vinylidene fluoride **copolymer** 12017-96-8, Chromium lithium oxide (CrLiO<sub>2</sub>) 12031-65-1, Lithium nickel oxide (LiNiO<sub>2</sub>) 12057-17-9, Lithium manganese oxide (LiMn<sub>2</sub>O<sub>4</sub>) 12162-79-7, Lithium manganese oxide limno<sub>2</sub> 12190-79-3, Cobalt lithium oxide (CoLiO<sub>2</sub>) 14283-07-9, Lithium tetrafluoroborate 21324-40-3, Lithium hexafluorophosphate 24937-79-9, **Pvdf** 25014-41-9, **Polyacrylonitrile** 25322-68-3, Peo 29935-35-1, Lithium hexafluoroarsenate 33454-82-9, Lithium triflate 90076-65-6 106818-19-3D, Ethylene carbonate-ethylene oxide copolymer, halogenated 131651-65-5 132843-44-8  
RL: DEV (Device component use); USES (Uses)  
(**secondary** battery using **porous film** type solvent-free polymer **electrolyte** filled with

oligomer/prepolymer **electrolyte**)  
 IT 25608-11-1DP, chloride terminated  
 RL: DEV (Device component use); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
 (**secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)  
 IT 1344-28-1, Alumina, uses 7631-86-9, Silica, uses 13463-67-7, Titania, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (**secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)  
 IT 37220-89-6, Lithium aluminate  
 RL: MOA (Modifier or additive use); USES (Uses)  
 ( $\gamma$ -; **secondary** battery using porous film type solvent-free polymer electrolyte filled with oligomer/prepolymer electrolyte)

L25 ANSWER 3 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2005:982634 HCAPLUS Full-text  
 DOCUMENT NUMBER: 143:249370  
 TITLE: Manufacture of porous films with improved porosity and air permeability  
 INVENTOR(S): Emori, Hideyuki; Aburaya, Aki  
 PATENT ASSIGNEE(S): Nitto Denko Corp., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 11 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 2005239773	A	20050908	JP 2004-48177	20040224
				20040224

PRIORITY APPLN. INFO.: JP 2004-48177

AB The films, useful for nonaq. **electrolyte** battery separators, are manufactured by kneading polymer compns. containing polyolefins and plasticizers, cooling the kneaded products, stretching the resulting sheets in  $\geq 1$  direction, eluting the plasticizers from the stretched products into **first** solvents, treatment of the stretched films with ultrasonic wave in nonpenetrable **second** solvents, and drying the stretched products. Thus, a composition containing **polyethylene**, Norsorex (norbornene rubber), TPE 821 (thermoplastic elastomer), and paraffin oil was kneaded, cooled, hot-pressed, biaxially stretched, treated with decane for elution of the paraffin oil, treated with ultrasonic wave in water, dried, and heat-treated for crosslinking to give a porous film with porosity 35%, air permeability (JIS P 8117) 380 s/100 cm<sup>3</sup>, and piercing strength 3.0 N.  
 IC ICM C08J009-26  
 ICS H01M002-16; C08L023-00  
 CC 38-2 (Plastics Fabrication and Uses)  
 Section cross-reference(s): 52  
 IT Drying  
 Plastic films  
 Plasticizers  
**Secondary** battery separators

(manufacture of porous films with improved porosity and air permeability)

IT **Polymer blends**

RL: TEM (Technical or engineered material use); USES (Uses)  
(manufacture of **porous films** with improved porosity and air permeability)

L25 ANSWER 4 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:611324 HCAPLUS Full-text

DOCUMENT NUMBER: 143:100453

TITLE: Separators for electronic devices

INVENTOR(S): Sugiyama, Masahide; Totsuka, Hiroki; Takahata, Masanori

PATENT ASSIGNEE(S): Tomoegawa Paper Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 13 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 2005190736	A	20050714	JP 2003-428047	20031224
			JP 2003-428047	20031224

PRIORITY APPLN. INFO.:

AB The separator comprises a **porous** elec. insulating **polymer film** containing polymers with **polyethylene** oxide structures and/or polymers with **polypropylene** oxide structures. Preferably, the separators contain Li salts. Such separators for **secondary** batteries, including (1) vinylidene fluoride homopolymer, (2) copolymer of vinylidene fluoride with tetrafluoroethylene, hexafluoropropylene, ethylene, propylene, and/or ethylene trifluoride, or (3) mixts. of (1) and (2) are also claimed. The electronic devices may especially be Li ion **secondary** batteries, polymer lithium batteries, aluminum **electrolytic** capacitors, or elec. double layer capacitors. The separators show low charging characteristics and excellent runnability and are suitable for mass production

IC ICM H01M002-16

ICS H01G009-02; H01M010-40

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 47, 76

ST separator porous elec insulator film; lithium salt PEO porous film separator; elec double layer capacitor **polypropylene** oxide separator; **polyethylene** oxide porous film separator **electrolytic** capacitor; **polyvinylidene fluoride** polymer battery separator; **porous** elec insulating **polymer film** separator

IT **Primary** battery separators

**Secondary** battery separators

(porous dielec. film separators containing polyethylene oxide and/or polypropylene oxide for batteries and capacitors)

IT **Electrolytic** capacitors

(separators for; porous dielec. film separators containing **polyethylene** oxide and/or **polypropylene** oxide for batteries and capacitors)

IT **Polymer electrolytes**



(separators; **porous** dielec. **film** separators  
containing **polyethylene** oxide and/or **polypropylene**  
oxide for batteries and capacitors)

IT 9002-88-4, Polyethylene 9003-07-0, Polypropylene 9011-14-7,  
Poly(methyl methacrylate) 24937-79-9, Vinylidene fluoride  
**homopolymer** 25014-41-9, Polyacrylonitrile

RL: DEV (Device component use); TEM (Technical or engineered  
material use); USES (Uses)

(**porous** separator **film**; **porous**  
dielec. **film** separators containing polyethylene oxide  
and/or polypropylene oxide for batteries and capacitors)

L25 ANSWER 5 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:239069 HCAPLUS Full-text

DOCUMENT NUMBER: 142:298942

TITLE: Method for continuous production of functional  
**film** having **porous**  
**resin** sheet containing functional  
polymer

INVENTOR(S): Hiraoka, Hideki; Kubota, Kouzou; Yamaguchi,  
Takeo; Ooya, Nobuo; Harada, Hiroshi

PATENT ASSIGNEE(S): Toagosei Co., Ltd., Japan

SOURCE: PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005023921	A1	20050317	WO 2004-JP6389	20040512
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
JP 2005076012	A	20050324	JP 2003-311841	20030903
CA 2537795	A1	20050317	CA 2004-2537795	20040512
PRIORITY APPLN. INFO.:				JP 2003-311841 A 20030903
				WO 2004-JP6389 W 20040512

- AB A method comprises (i) a step wherein a porous resin sheet (a porous **polyethylene** sheet or the like) is continuously conveyed and impregnated with a polymer precursor having a functional group (2-acrylamide-2-methylpropanesulfonic acid or the like), to attach the polymer precursor to the porous resin sheet, (ii) a polymerization step wherein a **first** and a **second** resin films (both are a polyester film or the like) are continuously supplied to both the surfaces of the porous sheet having been impregnated with the above polymer precursor and containing the polymer precursor attached thereto, in such a manner that the resin films are contacted with the surfaces and the porous sheet is sandwiched between the two resin films, and then the polymer precursor is polymerized (by the irradiation with an active energy ray, and the like), (iii) a film releasing step, (iv) and a polymer removing step. Thus, a **polyethylene** porous film was continuously impregnated with a solution comprising 2-acrylamido-2-methylpropanesulfonic acid 90, N,N'-methylenebisacrylamide 10, Darocure 1173 2, a surfactant 2, and water 100 parts, each **polyethylene** terephthalate film with oxygen permeability 385 mL/m<sup>2</sup>·24 h·MPa was placed on both side of the precursor-attached porous film, irradiated with a mercury lamp, removed two **polyethylene** terephthalate films to give an **electrolyte** membrane with semitransparency, good appearance, proton conductivity 53 S/cm<sup>2</sup>, and methanol permeation flux 0.72 kg/m<sup>2</sup>·hour.
- IC ICM C08J009-40
- CC 38-2 (Plastics Fabrication and Uses)  
Section cross-reference(s): 52
- ST continuous prodn functional **film porous resin** sheet; porous film contg acrylamidomethylpropanesulfonic acid methylenebisacrylamide copolymer electrolyte membrane
- IT Porous materials  
(films; method for continuous production of functional **films** having **porous resin** sheet containing functional polymers)
- IT Polymer electrolytes  
(method for continuous production of functional **films** having **porous resin** sheet containing functional polymers)
- IT Polyimides, uses  
RL: IMF (Industrial manufacture); POF (Polymer in formulation); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(polyether-, laminate with porous substrate; method for continuous production of functional **films** having **porous resin** sheet containing functional polymers)
- IT Polyethers, uses  
RL: IMF (Industrial manufacture); POF (Polymer in formulation); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(polyimide-, laminate with porous substrate; method for continuous production of functional **films** having **porous resin** sheet containing functional polymers)
- IT Polyolefins  
RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or engineered material use); USES (Uses)  
(porous substrates; method for continuous production of functional **films** having **porous resin** sheet containing functional polymers)
- IT Films  
(porous; method for continuous production of functional **films** having **porous resin** sheet containing functional polymers)
- IT 69824-22-2P, 2-Acrylamido-2-methylpropanesulfonic acid-N,N'-methylenebisacrylamide copolymer

RL: IMF (Industrial manufacture); MOA (Modifier or additive use);  
 PREP (Preparation); USES (Uses)  
 (functional polymer; method for continuous production of functional  
**films** having **porous resin** sheet  
 containing functional polymers)

IT 26298-81-7P 26615-45-2P

RL: IMF (Industrial manufacture); POF (Polymer in formulation); PRP  
 (Properties); TEM (Technical or engineered material use); PREP  
 (Preparation); USES (Uses)  
 (laminate with porous substrate; method for continuous production of  
 functional **films** having **porous resin**  
 sheet containing functional polymers)

IT 9002-88-4, Polyethylene 603951-77-5, U-Pore UP 3025

RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical  
 or engineered material use); USES (Uses)  
 (porous substrate; method for continuous production of functional  
**films** having **porous resin** sheet  
 containing functional polymers)

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR  
 THIS RECORD. ALL CITATIONS AVAILABLE IN  
 THE RE FORMAT

L25 ANSWER 6 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:894110 HCAPLUS Full-text

DOCUMENT NUMBER: 142:97389

TITLE: Preparation method of polymer electrolyte  
 product for lithium polymer **secondary**  
 battery and preparation method for the battery  
 using the polymer electrolyte product

INVENTOR(S): Do, Chil Hun; Jin, Bong Su; Jin, Chang Su; Mun,  
 Seong In; Yoon, Mun Su

PATENT ASSIGNEE(S): Korea Electro Technology Research Institute, S.  
 Korea

SOURCE: Repub. Korean Kongkae Taeho Kongbo, No pp. given  
 CODEN: KRXXA7

DOCUMENT TYPE: Patent

LANGUAGE: Korean

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
-----				
KR 2001090406	A	20011018	KR 2000-15364	200003 25

PRIORITY APPLN. INFO.: KR 2000-15364

200003  
25

AB A method for preparing the polymer electrolyte product for a lithium polymer  
**secondary** polymer and a method for preparing the lithium polymer **secondary** polymer  
 using the polymer electrolyte product are provided, to prevent the leakage of  
 liquid electrolyte. The preparation method of a polymer electrolyte product  
 comprises the steps of: dissolving poly(vinyl chloride) as a matrix polymer into a  
**first** solvent, adding a plasticizer with high b.p., adding silica whose surface is  
 coated with an organic material, and mixing them; coating the mixture onto a  
 substrate uniformly and drying it to make a polymer electrolyte film that the  
**first** solvent is removed; dipping the polymer electrolyte film into a **second**  
 solvent that the plasticizer with high b.p. is dissolved and the matrix polymer is

not dissolved, to remove only the plasticizer, and thereby making a polymer film with many micropores; and dipping the polymer film into a liquid electrolyte containing lithium salts for allowing the liquid electrolyte to soak into the micropores, and thereby obtaining the polymer electrolyte product. Preferably the number mean mol. weight of poly(vinyl chloride) is 10,000-200,000; the particle size of silica is less than 1  $\mu\text{m}$ ; the organic material is silane; the **first** solvent is tetrahydrofuran; the plasticizer is di-Bu phthalate or dioctyl phthalate; and the **second** solvent is methanol or Et ether.

- IC ICM H01M010-40
- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
- Section cross-reference(s): 38
- ST **secondary** lithium battery polymer **electrolyte**  
silica **PVC** porous composite
- IT Membranes, nonbiological  
(composite; preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT Porous materials  
(films; preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT **Secondary** batteries  
(lithium; preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT Dissolution  
(of polymer and plasticizers; preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT **Films**  
(porous; preparation method of **polymer** electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT Plasticizers  
Polymer electrolytes  
(preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT 7439-93-2D, Lithium, salts  
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)  
(composite with **PVC**/coated silica; preparation method of polymer **electrolyte** product for lithium polymer **secondary** battery and preparation method of battery using polymer **electrolyte** product)
- IT 9002-86-2  
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)  
(d.p. 162-3247; preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT 17341-24-1D, **PVC** complexes, uses  
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)  
(preparation method of polymer **electrolyte** product for lithium polymer **secondary** battery and preparation method of

1 layer

- battery using polymer **electrolyte** product)
- IT 60-29-7, Diethyl ether, uses 67-56-1, Methanol, uses 109-99-9, Tetrahydrofuran, uses  
 RL: NUU (Other use, unclassified); USES (Uses)  
 (preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT 84-74-2 117-81-7, Dioctyl phthalate  
 RL: PEP (Physical, engineering or chemical process); PYP (Physical process); REM (Removal or disposal); PROC (Process)  
 (preparation method of polymer electrolyte product for lithium polymer **secondary** battery and preparation method of battery using polymer electrolyte product)
- IT 7631-86-9, Silica, uses  
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)  
 (silane-coated, composite with PVC/Lithium salt complexes; preparation method of polymer **electrolyte** product for lithium polymer **secondary** battery and preparation method of battery using polymer **electrolyte** product)
- IT 7803-62-5, Silane, uses  
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)  
 (silica coated with, composite with PVC/lithium salt complexes; preparation method of polymer **electrolyte** product for lithium polymer **secondary** battery and preparation method of battery using polymer **electrolyte** product)

L25 ANSWER 7 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:889873 HCAPLUS Full-text

DOCUMENT NUMBER: 142:95120

TITLE: Polymeric electrolyte for electrochemical device using multilayered polymeric film as separator

INVENTOR(S): Ahn, Sun Ho; Hwang, Ji Yeong; Lee, Hyang Mok; Lee, Seung Jin

PATENT ASSIGNEE(S): LG Chem Investment Ltd., S. Korea

SOURCE: Repub. Korean Kongkae Taeho Kongbo, No pp. given  
 CODEN: KRXXA7

DOCUMENT TYPE: Patent

LANGUAGE: Korean

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
KR 2001055968	A	20010704	KR 1999-57312	19991213
PRIORITY APPLN. INFO.:				19991213
				19991213

AB Provided is a multilayered polymeric film having electrochem. stability, adhesive property with an electrode, **electrolytic** solution impregnating and stabilizing property, which is produced without extracting or removing plasticizer. And, a method for preparing thereof and a polymeric **electrolyte** for an electrochem.

device using the multilayered polymeric film as a separator are also provided. The multilayered polymeric film comprises: (i) a porous first polymer layer; and (ii) a gelled second polymer layer of polyvinylidene fluoride-chlorotrifluoroethylene copolymer. The method comprises the steps of: (i) dissolving polyvinylidene fluoride-chlorotrifluoroethylene copolymer in solvent selected from the group consisting of acetone, dimethylacetamide and N-methyl-2-pyrrolidone; and (ii) after coating or impregnating or simultaneously coating and impregnating one side or both sides of a porous polymer film with the solution of the step (i), drying it to form a gelled second polymer layer. The polymer electrolyte system for an electrochem. device comprises: (i) a multilayered polymeric separator comprising a porous first polymeric layer and a gelled second polymer layer of polyvinylidene fluoride-chlorotrifluoroethylene copolymer; and (ii) a liquid electrolyte comprising a salt and an organic solvent.

IC ICM B32B027-28

CC 38-3 (Plastics Fabrication and Uses)

Section cross-reference(s): 72, 76

L25 ANSWER 8 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2003:414438 HCAPLUS Full-text

DOCUMENT NUMBER: 138:404361

TITLE: Secondary nonaqueous electrolyte battery and its manufacture

INVENTOR(S): Segawa, Masazumi

PATENT ASSIGNEE(S): Japan Storage Battery Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 11 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 2003157898	A	20030530	JP 2001-355042	20011120

PRIORITY APPLN. INFO.:

JP 2001-355042

20011120

AB The battery has an anode and/or a cathode containing a 1st porous polymer electrolyte, and a 2nd porous polymer electrolyte layer (a) between the 2 electrodes or (b) on both sides of a separator; and is obtained by adhering (a) the electrode-electrolyte or (b) the cathode-electrolyte, separator-electrolyte and anode-electrolyte with heat treatment; where the battery satisfies  $(T2-4) \leq T1 \leq (T2+5)$  and  $T3 \geq (T2+10)$ , [ $T1$  = heat treatment temperature;  $T2$  = initial m.p. of the polymer electrolyte layer observed by DSC; and  $T3$  = reaction peak temperature].

IC ICM H01M010-40

ICS H01M004-02

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

ST porous polymer electrolyte secondary battery manuf; heat treatment DSC specification

IT Secondary batteries

(lithium; manufacture of electrodes containing porous polymer electrolytes for secondary lithium batteries)

IT Battery electrolytes

(manufacture of electrodes containing porous polymer electrolytes for secondary lithium batteries)

IT Fluoropolymers, uses

RL: DEV (Device component use); USES (Uses)  
 (manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 7440-50-8, Copper, uses  
 RL: DEV (Device component use); USES (Uses)  
 (anode collector; manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 7782-42-5, Graphite, uses  
 RL: DEV (Device component use); USES (Uses)  
 (anode; manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 7429-90-5, Aluminum, uses  
 RL: DEV (Device component use); USES (Uses)  
 (cathode collector; manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 143623-51-2, Cobalt lithium nickel oxide (Co<sub>0.15</sub>LiNi<sub>0.85</sub>O<sub>2</sub>)  
 RL: DEV (Device component use); USES (Uses)  
 (cathode; manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate  
 RL: DEV (Device component use); USES (Uses)  
 (electrolyte solution; manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 21324-40-3, Lithium hexafluorophosphate  
 RL: DEV (Device component use); USES (Uses)  
 (electrolyte; manufacture of electrodes containing porous polymer electrolytes for **secondary** lithium batteries)

IT 9011-17-0, Hexafluoropropylene-vinylidene fluoride copolymer  
 24937-79-9, **PVDF**  
 RL: DEV (Device component use); USES (Uses)  
 (manufacture of electrodes containing porous polymer **electrolytes** for **secondary** lithium batteries)

L25 ANSWER 9 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2003:71260 HCAPLUS Full-text

DOCUMENT NUMBER: 138:138410

TITLE: Porous polyolefinic films containing particles and production method thereof

INVENTOR(S): Kishii, Yutaka; Kii, Keisuke; Uetani, Yoshihiro; Fujita, Shigeru; Yamaguchi, Mutsuko

PATENT ASSIGNEE(S): Nitto Denko Corp., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 2003026847	A	20030129	JP 2001-214619	20010716
PRIORITY APPLN. INFO.:			JP 2001-214619	20010716

AB Title films useful for separators for nonaq. **electrolyte** batteries comprise polyolefin resins with weight average mol. weight  $\geq 5 + 105$  and particles with

particle diameter 0.001 - 10  $\mu\text{m}$ , and have surface pore size 0.01-1  $\mu\text{m}$ , pore size of the surface fractured in the thickness direction 0.1 - 10  $\mu\text{m}$ , and pore size ratio of surface/surface fractured in the thickness direction < 0.2. Thus, GUR 4012 (**polyethylene**) with m.p. 135° and weight average mol. weight 1 + 106 14, MG 35 (styrene-acrylate copolymer particle) with particle diameter 1  $\mu\text{m}$  8, and liquid paraffin 78 parts were kneaded at 160° and extruded at 160° to give a sheet-shaped article, which was preheated at 115°, a solvent was removed at 120°, stretched 2-folds in the width and longitudinal direction resp. to give a 14  $\mu\text{m}$ -thick porous film with porosity 53%, ion flow velocity 0.38 mmol/min, **electrolyte** impregnation ratio 32%, polymeric **electrolyte** impregnation ratio 43%, surface pore size 0.05  $\mu\text{m}$ , pore size of the surface fractured in the thickness direction 4.42  $\mu\text{m}$ , and pore size ratio of surface/surface fractured in the thickness direction 0.01, which was used as separator for a nonelectrolyte cell showing discharge capacity 1.42 mAh.

IC ICM C08J009-26  
ICS C08J009-28; C08K003-00; C08L023-06; C08L101-00; H01M002-16;  
H01M010-40  
CC 38-3 (Plastics Fabrication and Uses)  
Section cross-reference(s): 52, 76  
ST porous polyolefin film contg particle prepn; polyethylene styrene  
acrylate **copolymer resin particle porous**  
film prepn  
IT **Primary** batteries  
    **Secondary** batteries  
        (nonaq. electrolyte; preparation of porous polyolefinic films containing  
        particles useful for battery separators)  
IT **Primary** battery separators  
    **Secondary** battery separators  
        (nonaq.; preparation of porous polyolefinic films containing particles  
        useful for battery separators)  
IT **Polymer** electrolytes  
    (preparation of **porous polyolefinic films** containing  
    particles useful for battery separators containing)  
IT 79-10-7D, Acrylic acid, ester, polymers with styrene 100-42-5D,  
Styrene, polymers with acrylates  
RL: MOA (Modifier or additive use); USES (Uses)  
    (**resin particle**; preparation of **porous**  
    polyolefinic **films** containing particles useful for battery  
    separators)

L25 ANSWER 10 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN  
ACCESSION NUMBER: 2002:812223 HCAPLUS Full-text  
DOCUMENT NUMBER: 137:339975  
TITLE: Nonaqueous electrolyte **secondary**  
battery having **porous polymer**  
**layer** on depolarization **layer**  
and manufacture thereof  
INVENTOR(S): Tagawa, Masahiro; Kitano, Shinya; Hasumi,  
Takeshi  
PATENT ASSIGNEE(S): Japan Storage Battery Co., Ltd., Japan  
SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.  
CODEN: JKXXAF  
DOCUMENT TYPE: Patent  
LANGUAGE: Japanese  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	-----



JP 2002313429

A

20021025

JP 2001-120516

200104  
19

PRIORITY APPLN. INFO.:

JP 2001-120516

200104  
19

AB The nonaq. electrolyte **secondary** battery has a depolarization layer on a pos. electrode and/or a neg. electrode which has a porous polymer on the surface and in the pores, wherein a weight of the porous polymer per unit volume on the surface is higher than that in the pores and a thickness of the porous polymer on the surface is set at 0.1-6  $\mu\text{m}$ . The process comprises the steps of (1) applying an active agent on a metal foil current collector to form an electrode, (2) pressing the electrode, (3) impregnating the electrode with a polymer solution which has a viscosity  $\geq 1,000$  cps and contains a **1st** solvent, (4) extracting the **1st** solvent in the polymer on the electrode using a **2nd** solvent, and (5) drying the electrode.

IC ICM H01M010-40  
ICS H01M004-02; H01M004-04; H01M004-62

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 38

ST **porous polymer** depolarization layer  
nonaq electrolyte **secondary** battery

IT **Secondary** batteries  
(**porous polymer** in depolarization  
layer of nonaq. electrolyte **secondary** battery)

IT Polyoxyalkylenes, uses  
RL: DEV (Device component use); EPR (Engineering process); PEP  
(Physical, engineering or chemical process); PROC (Process); USES  
(Uses)  
(**porous polymer** in depolarization  
layer of nonaq. electrolyte **secondary** battery)

IT 9011-14-7, PMMA 9011-17-0, Hexafluoropropylene-vinylidene fluoride  
**copolymer** 25014-41-9, **Polyacrylonitrile**  
25322-68-3, PEO  
RL: DEV (Device component use); EPR (Engineering process); PEP  
(Physical, engineering or chemical process); PROC (Process); USES  
(Uses)  
(**porous polymer** in depolarization  
layer of nonaq. electrolyte **secondary**  
battery)

L25 ANSWER 11 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:672608 HCAPLUS Full-text

DOCUMENT NUMBER: 137:202356

TITLE: Ion-conducting adhesive **porous**  
**films, polymer** gelelectrolytes from them, their manufacture, and  
applicationsINVENTOR(S): Yamaguchi, Mutsuko; Uetani, Yoshihiro; Kii,  
Keisuke; Yamamura, Takashi; Nakamura, Seiji;  
Tabuchi, Masato

PATENT ASSIGNEE(S): Nitto Denko Corp., Japan; Daiso Co., Ltd.

SOURCE: Jpn. Kokai Tokkyo Koho, 20 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2002249742	A	20020906	JP 2001-358853	20011126
JP 2007035646	A	20070208	JP 2006-237184	20060901
PRIORITY APPLN. INFO.:			JP 2000-373466	A
			JP 2001-358853	A3
				20011126

- AB The films, showing 180° peeling strength  $\geq 2$  N/20 mm, comprise **porous** base **films** and **polymers** having poly(meth)acrylate, poly(ethylene oxide), poly(propylene oxide), poly(ethylene oxide/propylene oxide), polyphosphazene, poly(vinyl ether), or polysiloxane main chains and oligo(alkylene oxide) side chains. Polymer gel **electrolytes** manufactured using them are useful for batteries and capacitors. Thus, porous ultrahigh-mol.-weight **polyethylene** film was coated with a composition containing glycidyl methoxyethoxyethyl ether-allyl glycidyl ether-ethylene oxide (49:51:1) copolymer and Blemmer PDE 100, soaked in a solution containing LiClO<sub>4</sub>, and heated to give a gel showing conductivity  $8.0 + 10^{-4}$  S/cm.
- IC ICM C09J007-02  
ICS C08J009-36; C09J009-02; C09J171-00; C09J183-12; C09J201-00;  
H01B001-06; H01B013-00; H01G009-02; H01G009-035; H01G009-038;  
H01M010-40; C08L101-00
- CC 38-3 (Plastics Fabrication and Uses)  
Section cross-reference(s): 52, 76
- IT Porous materials  
(**films**; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT Capacitors  
Electrolytes  
Primary batteries  
Secondary batteries  
(ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT Polyethers, uses  
RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(oligo(alkylene oxide) side chain-containing; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT Polyoxyalkylenes, uses  
Polyphosphazenes  
Polysiloxanes, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(oligo(alkylene oxide) side chain-containing; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT Fluoropolymers, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(porous base **film**; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT Films

- (porous; ion-conducting adhesive **porous films**  
for **polymer** gel electrolytes)
- IT 7791-03-9, Lithium perchlorate  
RL: TEM (Technical or engineered material use); USES (Uses)  
(electrolyte; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT 115383-11-4P  
RL: IMF (Industrial manufacture); RCT (Reactant); PREP  
(Preparation); RACT (Reactant or reagent)  
(ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT 454171-46-1P  
RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT 9002-84-0, **PTFE**  
RL: TEM (Technical or engineered material use); USES (Uses)  
(porous base **film**; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)
- IT 9002-88-4, **Polyethylene**  
RL: TEM (Technical or engineered material use); USES (Uses)  
(ultrahigh-mol.-weight, porous base **film**; ion-conducting adhesive **porous films** for **polymer** gel electrolytes)

L25 ANSWER 12 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:499574 HCAPLUS Full-text

DOCUMENT NUMBER: 137:35458

TITLE: Crosslinked or modified **polymeric porous films** as separators for

INVENTOR(S): Kenichiro, Kami; Hiroshi, Ueshima; Ryuichirou, Shinkai; Norikazu, Hosokawa; Manabu, Yamada; Hideo, Amaki; Tomoaki, Tamura

PATENT ASSIGNEE(S): Denso Corp., Japan  
SOURCE: Fr. Demande, 75 pp.

CODEN: FRXXBL

DOCUMENT TYPE: Patent

LANGUAGE: French

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO. -----	KIND ----	DATE -----	APPLICATION NO. -----	DATE
FR 2814284	A1	20020322	FR 2001-12110	200109 19
JP 2002170541	A	20020614	JP 2001-145341	200105 15
PRIORITY APPLN. INFO.:			JP 2000-287145	A 200009 21
			JP 2001-145341	A 200105 15

JP 2000-141476

A

200005

15

- AB Non-swelling polymeric separators for batteries with non-aqueous **electrolytes** are prepared by crosslinking or modifying a **porous polymer film** with compds. that formed the crosslinks or added chains by at least 2 carbon atoms away from the linking units between the monomers (e.g., the ester linkage of polyesters). The polymeric film can be selected from polybenzimidazoles, **polyimides**, polyether-**polyimides**, polyamide-**polyimides**, polyphenylene sulfides, polyphenylene oxides, polyether-**polysulfones**, **polysulfones**, polyether-polyketones, aramides, saturated polyesters, polyoxymethylenes, etc. Suitable crosslinking agents or modifiers include acrylates, vinyl compds., and functionalized (unsatd.) alkoxysilanes.
- IC ICM H01M010-38
- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 38
- IT Polysiloxanes, uses  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
(acrylic, crosslinking or modifying agent; **polymeric porous films** as separators for batteries with nonaq. electrolytes)
- IT Polyamide fibers, uses  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
(aramid, crosslinked or modified, battery separators; **polymeric porous films** as separators for batteries with nonaq. electrolytes)
- IT Polyesters, uses  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
(aromatic, Vylon KS001, crosslinked or modified, battery separators; **polymeric porous films** as separators for batteries with nonaq. electrolytes)
- IT **Primary** battery separators  
**Secondary** battery separators  
(crosslinked or modified **polymeric porous films** as separators for batteries with nonaq. electrolytes)
- IT Fluoropolymers, uses  
Polyamides, uses  
Polybenzimidazoles  
Polyesters, uses  
**Polyimides**, uses  
Polyoxymethylenes, uses  
Polyoxyphenylenes  
**Polysulfones**, uses  
Polythiophenylenes  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
(crosslinked or modified, battery separators; **polymeric porous films** as separators for batteries with nonaq. electrolytes)
- IT Battery electrolytes  
(nonaq.; crosslinked or modified **polymeric porous films** as separators for batteries with nonaq. electrolytes)
- IT **Polyimides**, uses  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or

- reagent); USES (Uses)  
 (polyamide-, crosslinked or modified, battery separators;  
**polymeric porous films** as separators  
 for batteries with nonaq. **electrolytes**)
- IT **Polyimides**, uses  
 Polyketones  
**Polysulfones**, uses  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (polyether-, crosslinked or modified, battery separators;  
**polymeric porous films** as separators  
 for batteries with nonaq. **electrolytes**)
- IT Polyamides, uses  
 Polyethers, uses  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (polyimide-, crosslinked or modified, battery separators; **polymeric porous films**  
 as separators for batteries with nonaq. **electrolytes**)
- IT Polyethers, uses  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (polyketone-, crosslinked or modified, battery separators;  
**polymeric porous films** as separators  
 for batteries with nonaq. **electrolytes**)
- IT Acrylic polymers, uses  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (polysiloxane-, crosslinking or modifying agent;  
**polymeric porous films** as separators  
 for batteries with nonaq. **electrolytes**)
- IT Polyethers, uses  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (polysulfone-, crosslinked or modified, battery separators; **polymeric porous films**  
 as separators for batteries with nonaq. **electrolytes**)
- IT 2530-85-0,  $\gamma$ -(Methacryloxypropyl)trimethoxysilane  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (KBM503, crosslinking or modifying agent; **polymeric porous films** as separators for batteries with  
 nonaq. **electrolytes**)
- IT 4369-14-6, 2-Propenoic acid, 3-(trimethoxysilyl)propyl ester  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (KBM5103, crosslinking or modifying agent; **polymeric porous films** as separators for batteries with  
 nonaq. **electrolytes**)
- IT 1025-15-6, Triallyl isocyanurate  
 RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)  
 (TAIC, crosslinking or modifying agent; **polymeric porous films** as separators for batteries with  
 nonaq. **electrolytes**)
- IT 9016-80-2, Poly(methylpentene) 9020-73-9, **Polyethylene**  
 naphthalate 24937-79-9, **Poly(vinylidene fluoride)** 24968-11-4, **Polyethylene** naphthalate  
 24968-12-5, Polybutylene terephthalate 25038-59-9,  
**Polyethylene** terephthalate, uses 26062-94-2, Polybutylene

terephthalate 28779-82-0, Polybutylene naphthalate 51806-50-9,  
Polybutylene naphthalate  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or  
reagent); USES (Uses)

(crosslinked or modified, battery separators; **polymeric  
porous films** as separators for batteries with  
nonaq. **electrolytes**)

IT 75-94-5, Vinyltrichlorosilane 78-08-0, Vinyltriethoxysilane  
97-90-5, Ethylene glycol dimethacrylate 131-17-9, Diallyl  
phthalate 1067-53-4, Vinyltris( $\beta$ -methoxyethoxy)silane  
1321-74-0, Divinylbenzene, uses 1337-81-1, Vinylpyridine  
2768-02-7, Vinyltrimethoxysilane 3030-60-2, Allyl isocyanurate  
3290-92-4, Trimethylolpropane trimethacrylate 6294-79-7, Diallyl  
isocyanurate 21142-29-0,  $\gamma$ -(Methacryloxypropyl)triethoxysila  
ne 25013-15-4, Vinyltoluene 65100-04-1,  $\gamma$ -  
(Methacryloxypropyl)methyldiethoxysilane  
RL: DEV (Device component use); RCT (Reactant); RACT (Reactant or  
reagent); USES (Uses)  
(crosslinking or modifying agent; **polymeric  
porous films** as separators for batteries with  
nonaq. **electrolytes**)

L25 ANSWER 13 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:143078 HCAPLUS Full-text

DOCUMENT NUMBER: 136:201311

TITLE: Multicomponent composite film and method for  
preparing the same

INVENTOR(S): Lee, Seung-Jin; Lee, Hyang-Mok; Ahn, Soon-Ho;  
Cho, Jin-Yeon; Yong, Hyun-Hang; Lee, Hyung-Keun;  
Lee, Sang-Young; Song, Heon-Sik; Park,  
Soon-Yong; Kyung, You-Jin; Ahn, Byeong-In

PATENT ASSIGNEE(S): LG Chemical Co., Ltd., S. Korea

SOURCE: PCT Int. Appl., 34 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002015299	A1	20020221	WO 2001-KR1374	200108 11
W: CN, JP, US				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR				
KR 2002071203	A	20020912	KR 2001-11191	200103 05
EP 1310005	A1	20030514	EP 2001-958588	200108 11
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY, TR				
JP 2004506542	T	20040304	JP 2002-520328	200108 11

JP 3885100 B2 20070221  
 US 2002187401 A1 20021212 US 2002-110047  
 JP 2006289985 A 20061026 JP 2006-135816  
 20020405  
 20060515  
 PRIORITY APPLN. INFO.: KR 2000-46735 A 20000812  
 KR 2001-11191 A 20010305  
 JP 2002-520328 A3 20010811  
 WO 2001-KR1374 W 20010811

AB The present invention provides a multi-component composite film comprising (a) polymer support layer (e.g., **polypropylene**); and (b) **porous** gelable **polymer layer** (e.g., **polyvinylidene fluoride**) which is formed on one side or both sides of the support layer of (a), wherein the support film of (a) and the gelable polymer layer of (b) are unified without the interface, a method for preparing the same, and a polymer **electrolyte** system applied the same.

IC ICM H01M002-16

CC 38-3 (Plastics Fabrication and Uses)  
 Section cross-reference(s): 52

IT Fuel cells  
 Polymer electrolytes  
**Primary** batteries  
**Secondary** batteries  
 Sensors

(multicomponent composite film and method for preparing the same)

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR  
 THIS RECORD. ALL CITATIONS AVAILABLE IN  
 THE RE FORMAT

L25 ANSWER 14 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2000:420559 HCAPLUS Full-text

DOCUMENT NUMBER: 133:32683

TITLE: Fuel cell electrodes and their manufacture

INVENTOR(S): Hitomi, Shuji

PATENT ASSIGNEE(S): Japan Storage Battery Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 2000173624	A	20000623	JP 1999-78885	199903

DE 10004955	A1	20000817	DE 2000-10004955	24
				200002
				04
US 7147957	B1	20061212	US 2000-497515	200002
				04
PRIORITY APPLN. INFO.:		JP 1998-296157	A	199810
				03
		JP 1999-29045	A	199902
				05
		JP 1999-78885	A	199903
				24
		JP 1999-78889	A	199903
				24

AB The electrodes have a catalyst **layer** containing a solid **polymer** electrolyte on a **porous** conductive substrate, where the substrate contains a porous resin. The electrodes are prepared by: impregnating a porous conductive material with a solution of a resin dissolved in a **1st** solvent, immersing the impregnated material in a **2nd** solvent miscible with the **1st** solvent but insol. for the **resin** to form a **porous resin layer** on the material, and joining the material with an electrolyte containing catalyst layer.

IC ICM H01M004-86  
ICS H01M004-88; H01M008-02; H01M008-10

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT Fuel cell electrodes  
(structure and manufacture of fuel cell electrodes containing polymer electrolyte-catalyst **layer** on **porous polymer** coated carbon substrates)

IT Fluoropolymers, uses  
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
(structure and manufacture of fuel cell electrodes containing polymer electrolyte-catalyst **layer** on **porous polymer** coated carbon substrates)

IT 7440-06-4, Platinum, uses  
RL: CAT (Catalyst use); USES (Uses)  
(structure and manufacture of fuel cell electrodes containing polymer electrolyte-catalyst **layer** on **porous polymer** coated carbon substrates)

IT 7440-44-0, Carbon, uses 24937-79-9, Poly(**vinylidene fluoride**)  
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
(structure and manufacture of fuel cell electrodes containing polymer electrolyte-catalyst **layer** on **porous polymer** coated carbon substrates)



TITLE: Porous polymer film  
 as separator for nonaqueous-electrolyte battery  
 INVENTOR(S): Ishibashi, Mamoru  
 PATENT ASSIGNEE(S): Asahi Chemical Industry Co., Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 8 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 11086830	A	19990330	JP 1997-241098	199709 05
PRIORITY APPLN. INFO.: JP 1997-241098				199709 05

AB The title film comprises a polar polymer having continuous pores in the film and thickness direction, and the pores at either pair of the opposing end faces are closed. The film absorbs the electrolyte rapidly and has high leakage-preventing ability.

IC ICM H01M002-18  
 ICS H01M002-16; C08J009-28

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 38

ST polar polymer separator nonaq **electrolyte** battery; porous battery separator rapid **electrolyte** impregnation; leakage prevention porous battery separator; acrylonitrile polymer battery separator; **polyvinylidene fluoride** polymer battery separator

IT Porous materials  
 (films; porous polar polymer film as separator for nonaq.-electrolyte battery for rapid electrolyte impregnation and leakage prevention)

IT **Primary** battery separators  
**Secondary** battery separators  
 (porous polar polymer film as separator for nonaq.-electrolyte battery for rapid electrolyte impregnation and leakage prevention)

IT Fluoropolymers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (porous polar polymer film as separator for nonaq.-electrolyte battery for rapid electrolyte impregnation and leakage prevention)

IT **Films**  
 (porous; porous polar polymer film as separator for nonaq.-electrolyte battery for rapid electrolyte impregnation and leakage prevention)

IT 9011-17-0, Kynar 2801 24937-79-9, KF 1000 25014-41-9, Acrylonitrile **homopolymer** 26658-88-8, Acrylonitrile-methyl.acrylate-sodium methallylsulfonate **copolymer**  
 RL: DEV (Device component use); USES (Uses)  
 (porous polar polymer film as separator for nonaq.-electrolyte battery for rapid electrolyte impregnation and leakage prevention)

L25 ANSWER 16 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1997:61043 HCAPLUS Full-text

DOCUMENT NUMBER: 126:77478

TITLE: Nonaqueous electrolyte batteries with single sheet porous polymer separators

INVENTOR(S): Gotanda, Yukihiro; Terao, Yoshiki; Sato, Hiromi

PATENT ASSIGNEE(S): Matsushita Electric Ind Co Ltd, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 08293296	A	19961105	JP 1995-98224	19950424
PRIORITY APPLN. INFO.:				JP 1995-98224
				19950424

AB The batteries use separators composed of a single porous polymer sheet having 5-50  $\mu\text{m}$  thickness, with the (1-10)- $\mu\text{m}$  thick layer on both sides having porosity 1-10% and the remaining part having porosity 50-95%. The polymer is preferably a polyolefin. These batteries have good performance at high current.

IC ICM H01M002-16

ICS H01M006-16; H01M010-40

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT **Primary** battery separators**Secondary** battery separators(single sheet porous **polymer** separators with low **porosity** surface **layers** for nonaq. electrolyte batteries)IT 9003-07-0, **Polypropylene**

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(single sheet porous **polymer** separators with low **porosity** surface **layers** for nonaq. **electrolyte** batteries)

L25 ANSWER 17 OF 17 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:539616 HCAPLUS Full-text

DOCUMENT NUMBER: 115:139616

TITLE: Sealed lead-acid batteries

INVENTOR(S): Tokunaga, Akio

PATENT ASSIGNEE(S): Japan Storage Battery Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	-----

JP 03057166

A

19910312

JP 1989-192202

198907

25

PRIORITY APPLN. INFO.:

JP 1989-192202

198907

25

AB The batteries have spacers between electrodes in an electrode stack and an electrolyte retainer filled between the electrodes and around the stack. The spacers are corrugated acid- and oxidation-resistant synthetic **resin films** having >70% **perforation**, the electrolyte retainer is fine SiO<sub>2</sub> prepared from a **primary** powder having particle diameter 10-40  $\mu$ m, and the electrolyte is impregnated and retained in the retainer. These batteries have good performance and are easy to manufacture

IC ICM H01M010-10  
ICS H01M010-12

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT Batteries, **secondary**  
(sealed, lead-acid, with silica **electrolyte** retainers  
and corrugated perforated **polypropylene** spacers)

=> d 126 ibib abs hitstr hitind 1-15

L26 ANSWER 1 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2003:142811 HCAPLUS Full-text

DOCUMENT NUMBER: 139:244973

TITLE: Studies on modified atmosphere packaging on quality stability of intact or pre-cut welsh onion

AUTHOR(S): Ibaraki, Toshiyuki

CORPORATE SOURCE: Fukuoka Agric. Res. Cent., Chikushino, Japan

SOURCE: Fukuoka-ken Nogyo Sogo Shikenjo Tokubetsu Hokoku (2002), 18, 1-83

CODEN: FSSHEQ; ISSN: 0913-509X

PUBLISHER: Fukuoka-ken Nogyo Sogo Shikenjo

DOCUMENT TYPE: Journal

LANGUAGE: Japanese

AB Modified atmospheric packaging (MAP) which involves the lowering of oxygen (O<sub>2</sub>) and increasing of carbon dioxide (CO<sub>2</sub>) content maintains the quality of fresh vegetables and fruits caused by reduced respiration rates, weight loss, chemical component loss, and delay in yellowing. However, the gas composition surrounding fresh commodities is influenced by the respiration rate of the commodity and the gas permeability of the packaging film. Therefore, use of an improper packaging film may induce an off-odor as a result of anaerobic respiration. The objective of this study was to determine the effect of MAP on quality stability of intact or pre-cut Welsh onion (*Allium fistulosum* L.). The respiration rate, the chemical components including reduced ascorbic acid (RAA) and sugar, and the quality stability of Welsh onion harvested in summer (July, '95), autumn (Nov., '95), winter (Jan., '96) and spring (Apr., '96) were investigated during storage at room temperature or at 15°. A quadratic regression equation ( $Y = a + b + T + c + T^2$ ), with high relative coefficient ( $R = 0.993-0.995$ ), was established between respiration rate ( $\text{mgCO}_2 \cdot \text{kg}^{-1} \cdot \text{hr}^{-1}$ ) (Y) and temperature of Welsh onion (°C) (T). At the same temperature of 15°, Welsh onion harvested in winter season showed higher respiration rate than that harvested in summer season. In the Welsh onion harvested in summer and stored at room temperature condition, RAA content decreased markedly and the score of leaf tip withering increased. These indicate that the Welsh onion harvested in summer season and stored at room temperature does not retain good quality. RAA and sugar contents in Welsh onion harvested in winter were kept for 6 days, suggesting that it retains good quality during

storage at room temperature. On the other hand, summer season Welsh onion stored at 15° indicated much better quality stability than that of winter season. There was a correlation between the respiration rate of Welsh onion and quality stability, meaning that the lower was the respiration rate, the higher was the quality stability. Respiration rates in packages with various gas mixts. were measured by use of a closed system. An empirical equation for the respiration rate of Welsh onion was estimated as a function of the O<sub>2</sub> and CO<sub>2</sub> concentration. The equation that was obtained by multiple regression anal. has a significantly high correlation coefficient (0.92) and estimated as follows.  $R = 30.40 - 2.06 + [CO_2] + 0.57 + [O_2]$ . The effects of several atmospheric composition on respiration, nutrients and quality stability of Welsh onion were investigated during storage at 15°. Carbon dioxide production from Welsh onion under continuous stream of air (the flow rate was 7 liter·hr<sup>-1</sup>) was 111 mgCO<sub>2</sub>·kg<sup>-1</sup>·hr<sup>-1</sup>, it was reduced under continuous streams of low oxygen and high carbon dioxide levels. RAA, sugar and chlorophyll retention were better under low oxygen level than that in air. Sensory score of Welsh onion withering leaf tip was reduced under low oxygen levels. From these results, atmospheric compns. such as including 7.6% O<sub>2</sub> and 12.6% CO<sub>2</sub>, 4.1% O<sub>2</sub> and 17.1% CO<sub>2</sub> resp., gave better quality stability of Welsh onion while avoiding physiol. injury. Using a math. model employing previous equation of respiration rate, the gas composition inside the plastic film package and shipping container was predicted. A theor. model for atmospheric in a polymeric film package containing fresh commodities and in a shipping container was developed and validated by expts. with or without Welsh onion. The model was of three processes: respiration of fresh commodities, and **permeability of polymeric film** and of the shipping container. When vacant packages were used, changes in CO<sub>2</sub> and O<sub>2</sub> concns. inside the bi-axial oriented **polypropylene** (OPP) film package and expanded polystyrene (EPS) container agreed very well with simulated data. The fit indicates the suitability of the proposed model. Changes in O<sub>2</sub> and CO<sub>2</sub> concns. in the EPS container and O<sub>2</sub> concentration in the OPP film package were approximated when Welsh onion was packed in the film package. The predicted O<sub>2</sub> and CO<sub>2</sub> concentration in the film package differed slightly from the exptl. data with a maximal of 1.5% and 1.2%, resp. However, these exptl. and simulated gas concns. correlated significantly well, indicating that the simulation model can be useful for fresh commodities wrapped both with polymeric film and shipping container. This math. model, therefore, is useful for designing MAP system using polymeric film while in shipping containers with gas permeability. Assuming that CO<sub>2</sub>, O<sub>2</sub> and N<sub>2</sub> **permeability of polymeric film** is 4100, 1700 and 1300 (mL·m<sup>-2</sup>·day<sup>-1</sup>·atm<sup>-1</sup>), resp., O<sub>2</sub> concentration in polymeric film will be about 7-8% after 48 h. OPP film is widely used for packaging of fresh vegetables in Japan because of its transparency and adaptability for packaging machinery. Generally, the gas permeability of OPP film is too low to maintain the gas conditions suitable for vegetables. If fresh vegetables are wrapped with completely heat-sealed OPP film and shipped at high temperature, off-odors will result and the quality will deteriorate. So, chemical components and quality of Welsh onion wrapped roughly or completely heat-sealed with OPP film were investigated during storage at 5 and 20°. Concns. of O<sub>2</sub> and CO<sub>2</sub> in the roughly heat-sealed packages were the same as in air, whereas O<sub>2</sub> concentration in the completely heat-sealed packages decreased to 2-3% at 20° and to 4-5% at 5°. Contents of RAA and chlorophyll in onion which was roughly sealed and stored at 20° decreased, and withering of leaf tip occurred. This meant that the quality deteriorated severely. However, the contents of RAA and chlorophyll in Welsh onion is which were sealed completely and stored at 20°, and which were sealed roughly or completely and stored at 5°, did not change or only slightly decreased. In Welsh onion sealed completely and stored at 20°, off-odor was detected after 2 days and severely after 7 days and leaf tips began to wither after 7 days, suggesting that quality could be kept only for 4 days storage. In Welsh onion sealed completely and stored at 5°, off-odors and withering of leaf tip was not observed for 9 days, indicating that quality marketability could be kept for that period. The effects of combinations of OPP film packages and shipping containers on quality stability of Welsh onion were investigated under actual conditions. In

an actual transportation, there was only a small difference in the Welsh onion temperature between EPS container and usual corrugated fiberboard container. The result suggests that both materials have the almost same holding temperature effect. On the other hand, the gas permeability of OPP film is too low to maintain the gas conditions suitable for intact Welsh onion. Recently, OPP film with micro-perforations is developed owing to avoid the anaerobic respiration. Gas permeability of this film is determined by film itself and perforation. So adjusting the diameter or number of micro-perforations, gas permeability of this film could be suitable for this vegetable. In an actual transportation, O<sub>2</sub> concentration in the completely heat-sealed OPP film packages in the EPS container decreased to 2%. Therefore, the onion was induced to anaerobic respiration. O<sub>2</sub> concentration in completely heat-sealed OPP film packages with 5 micro-perforations in the corrugated fiberboard container decreased to 4.5% at the container was opened, thereafter increased to 7-9%. Therefore, this film made available to prevent the deterioration caused by anaerobic respiration. So the sugar and chlorophyll contents were comparatively retained. Too many micro-perforations were ineffective, however, because the O<sub>2</sub> concentration in the package was too high to maintain the MAP condition. To determine the method for evaluating the quality of pre-cut Welsh onion, quality of cut onion was examined and compared with elec. conductivity (EC) and potassium contents of **electrolyte** solution extracted from the onion. Quality of cut onions which were stored at 0, 5 and 10° were stable for 6 days. On the other hand, the quality of onions stored at 15 and 20° slightly deteriorated after 4 and 3 days of storage, resp., and thereafter becoming severe. Degree of EC of onions stored at 15 and 20° decreased for **first** 4 and 3 days of storage, resp., and thereafter began to increase, though degree of EC of onions stored at 0, 5 and 10° decreased during storage. The changes in potassium content extracted by homogenization was steady during storage, although the changes in potassium extracted by leakage similar to those in EC stored at 15 or 20°. Considering the relationship between degree of EC or potassium content and quality of cut onion, **electrolyte** leaked from openings when the tissue was collapsed due to decay. So the **electrolyte** leakage was good parameter to evaluate the decay of onion. Moreover, chlorophyll content is a useful parameter to evaluate the quality of cut onion. Chemical components and quality stability of pre-cut Welsh onion wrapped with unperforated or perforated OPP film package were investigated during storage at 10°. The concentration of O<sub>2</sub> in the unperforated OPP film package decreased rapidly and dropped to 2% while CO<sub>2</sub> increase. Therefore the onion was induced to anaerobic respiration. So the ethanol was produced and chemical components were reduced. Moreover the cellular of this onion became water-soaked. O<sub>2</sub> concentration in the OPP film package with too more micro-perforations decreased slightly. So ethanol was not detected and chemical components were decreased. Moreover the quality of this onion decreased and became unmarketable after 6 days of storage. In cut Welsh onion which was packed with perforated OPP film package (with 24 micro-perforations), O<sub>2</sub> concentration in this package was not considered so low as MAP for onion, but decreased and dropped to 10%. So ethanol could not be detected and chemical components could be kept or decreased slightly. Therefore the cut Welsh onion was acceptable for sale for 6 days of storage. These results suggested that MAP is effective technique for maintaining the quality of intact or pre-cut Welsh onion. A theor. model for atmospheric in a polymeric film package containing fresh commodities and in a shipping container was developed. The simulation model can be useful for Welsh onion wrapped both with OPP film and EPS container. The gas permeability of OPP film was too low to maintain the gas condition for intact or pre-cut Welsh onion; an OPP film with micro-perforations was available for avoiding the deterioration caused by anaerobic respiration. It appears the gas composition in the package can extend the shelf-life of the commodity. But too more micro-perforations was ineffective because of the O<sub>2</sub> concentration in the package could not be so low to maintain the MAP condition.

CC 17-10 (Food and Feed Chemistry)

ACCESSION NUMBER: 2001:713842 HCAPLUS Full-text  
 DOCUMENT NUMBER: 135:259856  
 TITLE: Method of fabrication of polymer electrolyte  
 battery of high mechanical strength and high  
 heat resistance  
 INVENTOR(S): Yamasaki, Mikiya  
 PATENT ASSIGNEE(S): Sanyo Electric Co., Ltd., Japan  
 SOURCE: U.S. Pat. Appl. Publ., 9 pp.  
 CODEN: USXXCO  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO. -----	KIND ----	DATE -----	APPLICATION NO. -----	DATE
US 2001024756	A1	20010927	US 2001-815009	200103 23
US 6623891	B2	20030923		
JP 2001273929	A	20011005	JP 2000-87627	200003 27
EP 1139479	A2	20011004	EP 2001-106718	200103 16
EP 1139479	A3	20050817		
EP 1139479	B1	20060726		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
TW 492208	B	20020621	TW 2001-90106569	200103 21
CA 2342368	A1	20010927	CA 2001-2342368	200103 26
CN 1316792	A	20011010	CN 2001-116500	200103 27
HK 1039405	A1	20070323	HK 2002-100896	200202 05
PRIORITY APPLN. INFO.:			JP 2000-87627	A 200003 27

AB A polymer electrolyte battery having a high discharge capacity, a high mech. strength, and a high heat resistance is provided. This polymer electrolyte battery includes: a polymer electrolyte layer covering an edge face of a pos. electrode plate and an edge part of a **porous** membrane; a **polymer** electrolyte **layer** covering an edge face of a neg. electrode plate and an edge part of the **porous** membrane; and a **polymer** electrolyte **layer** covering an edge face of the porous membrane in a manner that connects the polymer electrolyte layer with the polymer electrolyte layer.

IC ICM H01M010-40  
 ICS H01M006-16; H01M006-18

INCL 429303000

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 37

IT Battery electrolytes  
**Primary** batteries  
 Strength  
 Thermal resistance  
 (method of fabrication of polymer electrolyte battery of high mech. strength and high heat resistance)

IT 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate 7429-90-5, Aluminum, uses 7782-42-5, Graphite, uses 9002-88-4, **Polyethylene** 12190-79-3, cobalt lithium oxide colio2 21324-40-3, Lithium hexafluorophosphate  
 RL: DEV (Device component use); USES (Uses)  
 (method of fabrication of polymer **electrolyte** battery of high mech. strength and high heat resistance)

IT 9003-07-0, **Polypropylene** 52496-08-9, **Polypropylene** glycol diacrylate  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (method of fabrication of polymer **electrolyte** battery of high mech. strength and high heat resistance)

L26 ANSWER 3 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 1999:387727 HCAPLUS Full-text  
 DOCUMENT NUMBER: 131:21306  
 TITLE: Porous films, battery separators using the films, and batteries  
 INVENTOR(S): Matsushita, Kiichiro; Higuchi, Hiroyuki; Wano, Takashi; Ishisaki, Akira  
 PATENT ASSIGNEE(S): Nitto Denko Corp., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 11158304	A	19990615	JP 1998-234740	199808 20
PRIORITY APPLN. INFO.:			JP 1997-258950	A 199709 24

AB The films, when measured in an organic **electrolyte** solution at .apprx.20°, have a membrane resistance  $\leq 5 \Omega \cdot \text{cm}^2$ , which increased to  $\geq 1000 \Omega \cdot \text{cm}^2$  after hot pressing for 1 s at 100-130° and 5 kg/cm<sup>2</sup> for pore sealing.. Preferably, the films have a gas permeability  $\leq 400 \text{ s}/100 \text{ cm}^3$  and contain 10-80% of a **polyethylene**, having crystallinity  $\geq 50\%$  and weight average mo. weight 500-10,000, and 20-90% of a polymer having a m.p. higher by  $\geq 5^\circ$  than that of the **polyethylene**.

IC ICM C08J009-00  
 ICS H01M002-16; H01M006-16; H01M010-40; C08L023-04; C08L023-10

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

ST battery separator **porous polymer** blend  
**film**; polyethylene blend porous film separator characteristics

IT **Primary** battery separators  
 (characteristics of **porous** polyethylene containing **polymer** blend **films** for battery separators)

IT 9002-88-4, Polyethylene  
 RL: DEV (Device component use); PRP (Properties); USES (Uses)  
 (characteristics of **porous** polyethylene containing  
**polymer** blend **films** for battery separators)

L26 ANSWER 4 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1998:586414 HCAPLUS Full-text  
 DOCUMENT NUMBER: 129:262849  
 TITLE: Porous films and battery separators with  
 improved low-temperature shut-down capability  
 therefrom  
 INVENTOR(S): Wano, Takashi; Nishiyama, Souji; Matsushita,  
 Kiichiro  
 PATENT ASSIGNEE(S): Nitto Denko Corp., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 5 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 10237202	A	19980908	JP 1997-42710	199702 26

PRIORITY APPLN. INFO.: JP 1997-42710  
 199702  
 26

AB The title  $\geq 3$ -layer films, suitable for separators of nonaq. **electrolytic** solns. in batteries, consist of at least (a) a middle layer prepared from mixts. of **polyethylene** (I; melt index  $\leq 0.35$ ) and **polypropylene** (II) and (b) layers of II on the outsides of the middle layer and satisfy the relation  $2\% \leq \text{I content} < 30\%$ . Thus, isotactic II and an 80:20 mixture of HDPE (MI 0.3) and isotactic II were extruded to give a 3-layer film, which was heat-treated at  $135^\circ$  for 60 h, stretched, and shrunk. The resulting porous film showed I 20%, a peel strength of 100 g/10 mm, and a shut-down initiation temperature of  $126^\circ$ .

IC ICM C08J009-00

ICS B32B027-32; H01M002-16

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 38

IT **Primary** battery separators  
 (from **porous** multilayer **polymer** films  
 )

IT **Polymer** blends

RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical  
 or engineered material use); USES (Uses)  
 (**porous** multilayer **films** for battery  
 separators from)

L26 ANSWER 5 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1996:644057 HCAPLUS Full-text  
 DOCUMENT NUMBER: 125:280714  
 TITLE: Cylindrical air batteries  
 INVENTOR(S): Noya, Shigeto; Fujiwara, Takafumi; Watanabe,  
 Tomoya; Kubo, Isao; Kobayashi, Shigeo  
 PATENT ASSIGNEE(S): Matsushita Electric Ind Co Ltd, Japan



SOURCE: Jpn. Kokai Tokkyo Koho, 7 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 08222231	A	19960830	JP 1995-24128	199502 13
JP 3232936	B2	20011126		
JP 2002063948	A	20020228	JP 2001-205872	200107 06
JP 3642298	B2	20050427		
JP 3642297	B2	20050427	JP 2001-205871	200107 06

PRIORITY APPLN. INFO.: JP 1995-24128 A3  
 199502  
 13

AB The batteries use cathodes having a tubular catalyst layer, containing a **polytetrafluoroethylene** dispersed Ni film plated on a current collector, and a gas **permeable porous hydrophobic resin film** on the outside of the catalyst layer. The batteries have a tubular cathode with its lower end inserted in a ring shaped recession of a polymer member, which has a Sn plated brass ring on the inside and fit on a polymer sealing piece. The batteries have a tubular cathode with its open covered with an inside metal cap and an outside metal cap, a tubular separator with a sealed bottom inside the tubular cathode, an anode material free gelled **electrolyte** on the inside of the separator facing the inside of the inside cap, and a gelled Zn anode filled in the remaining part of the separator tube.

IC ICM H01M004-86

ICS H01M004-66; H01M012-06

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT Batteries, **primary**

(structure of cylindrical air batteries for preventing electrolyte leakage)

L26 ANSWER 6 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1990:500881 HCAPLUS Full-text

DOCUMENT NUMBER: 113:100881

TITLE: Batteries with laminated air-diffusion films

INVENTOR(S): Fukuda, Hiroshi; Yanagihara, Nobuyuki; Takada, Kanji; Yoshino, Masaaki

PATENT ASSIGNEE(S): Matsushita Electric Industrial Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 01267974	A	19891025	JP 1988-96005	

198804

19

PRIORITY APPLN. INFO.:

JP 1988-96005

198804

19

AB Batteries using air-diffusion cathodes have air-inlet openings on their cases and a vinyltrimethylsilane-hexamethylcyclotrisiloxane copolymer film supported with  $\geq 1$  porous film between the cases and the cathodes. Porous **polypropylene** films and nonwoven **polypropylene** fabrics were used as supports in examples. These films prevent **electrolyte** leakage from the batteries and increase their shelf life.

IC ICM H01M012-06

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

Section cross-reference(s): 38

IT Batteries, **primary**

(air-zinc, with air-diffusion films from vinyltrimethylsilane-hexamethylcyclotrisiloxane **copolymer** and **porous** supports)

L26 ANSWER 7 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1990:482132 HCAPLUS Full-text

DOCUMENT NUMBER: 113:82132

TITLE: Batteries with laminated air-diffusion films

INVENTOR(S): Fukuda, Hiroshi; Yanagihara, Nobuyuki; Takada, Kanji; Yoshino, Masaaki

PATENT ASSIGNEE(S): Matsushita Electric Industrial Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 01267972	A	19891025	JP 1988-96003	198804 19

PRIORITY APPLN. INFO.:

JP 1988-96003

198804

19

AB Batteries using air-diffusion cathodes have air inlet openings on their battery cases and a vinylphenol-di-Me siloxane block copolymer film supported with  $\geq 1$  porous films between the inside surface of the cases and the cathodes. Porous **polypropylene** films and nonwoven **polypropylene** fabrics were used as supports. These films prevent **electrolyte** leakage from the batteries and increase the shelf life of the batteries.

IC ICM H01M012-06

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT Batteries, **primary**

(air-zinc, air-diffusion films for, containing vinylphenol-di-Me siloxane **copolymer** films and **porous** supporting films)

L26 ANSWER 8 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1989:481388 HCAPLUS Full-text

DOCUMENT NUMBER: 111:81388

TITLE: Air battery

INVENTOR(S): Okazaki, Ryoji; Ito, Zenichiro; Takada, Kanji  
 PATENT ASSIGNEE(S): Matsushita Electric Industrial Co., Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 7 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 01059782	A	19890307	JP 1987-215647	19870828
				19870828

PRIORITY APPLN. INFO.: JP 1987-215647

AB The battery containing a gas-diffusion O cathode has a composite film of  $\geq 1$  microporous film supported by a thin polymethylpentene film inserted between the battery case and the cathode. The microporous film can be a polyolefin (**polypropylene** or **polyethylene**), a fluororesin, or a **polysulfone**, and is on the battery-case side of the composite. The polymethylpentene film can be in direct contact with the cathode or have a porous **PTFE** film in between. Air batteries using this composite do not show **electrolyte** leak and have good performance at high and low loads.

IC ICM H01M012-06

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT Batteries, **primary**  
 (air, polymethylpentene-microporous polymer composites in, for preventing electrolyte leak)

IT Fluoropolymers  
 RL: USES (Uses)  
 (films, composites of **polymethylpentene** and **microporous**, for air batteries)

IT 9016-80-2, Poly(methylpentene)  
 RL: USES (Uses)  
 (composites of **microporous polymer films** and, for air batteries)

IT 9002-88-4, Polyethylene 9003-07-0, Polypropylene  
 RL: USES (Uses)  
 (films, composites of **polymethylpentene** and **microporous**, for air batteries)

L26 ANSWER 9 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 1988:634174 HCAPLUS Full-text  
 DOCUMENT NUMBER: 109:234174  
 TITLE: Alkaline batteries with porous separators  
 INVENTOR(S): Ikeda, Konosuke  
 PATENT ASSIGNEE(S): Sanyo Duracell Co., Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

JP 63158741

A

19880701

JP 1986-305021

198612

19

PRIORITY APPLN. INFO.:

JP 1986-305021

198612

19

AB Alkaline Zn-MnO<sub>2</sub> batteries have **porous synthetic-resin film** separators with or without alkaline-resistant nonwoven cloth of natural or synthetic fibers on the anode side. The film is **polyethylene** film; and the cloth is made of **nylon**, **vinylon**, **polypropylene**, or cotton fibers. Thus, when discharged through a 3.9- $\Omega$  resistor to 0.9-V cutoff, Zn-MnO<sub>2</sub> batteries using 20- $\mu$ m porous **polyethylene** separators and a ZnO-saturated 40% KOH **electrolyte** had a discharge durations of 4.30 and 3.40 h before and after a 30-day storage at 60° vs. 4.44 and 3.46 h for batteries using a 0.1-mm-thick nonwoven vinylon separator. No leak was observed on batteries of the invention after storage, however 20% of the latter batteries showed leak.

IC ICM H01M002-16

ICS H01M006-06

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 38

IT Batteries, **primary**  
(separators, porous polyethylene films)

L26 ANSWER 10 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1988:613606 HCAPLUS Full-text

DOCUMENT NUMBER: 109:213606

TITLE: Lithium batteries with metal-coated separators

INVENTOR(S): Imaizumi, Masahiko; Niso, Kiyoshi; Sasama,  
Hiroshi; Okamoto, Osamu; Iwamaru, Futayasu

PATENT ASSIGNEE(S): Hitachi Maxell, Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	
JP 63143744	A	19880616	JP 1986-290920	198612 06

PRIORITY APPLN. INFO.:

JP 1986-290920

198612

06

AB **Microporous resin films** for use as separators in Li batteries have a metal electrochem. alloyed with Li on its anode-facing side. The metal is selected from Al, Sn, Zn, Pb, Bi, Si, Sb, and/or Mg. Thus, a 5- $\mu$ m Al layer was vapor deposited on a microporous **polypropylene** separator for use in a button-type Li-MnO<sub>2</sub> battery with a 1M LiClO<sub>4</sub>/2:1 (volume) propylene carbonate-MeOC<sub>2</sub>H<sub>4</sub>OMe **electrolyte**. The Al layer was electrochem. alloyed with Li prior to assembling into the battery. Batteries of the invention had internal resistances of 10-15  $\Omega$  and closed-circuit voltages of 2.45-2.52 V at the end of a 0.2-s discharge through a 560- $\Omega$  load at 10° after a 160-h predischarging through

an 18-k $\Omega$  load at 25° vs. 15-20  $\Omega$  and 2.30-2.42 V for batteries using a sep. 5- $\mu$ m Al sheet between the anode and uncoated separator.

IC ICM H01M004-06  
ICS H01M004-02; H01M004-40  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 38  
IT Batteries, **primary**  
(separators, polypropylene, with aluminum-lithium alloy layers,  
for low resistance)

L26 ANSWER 11 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1988:593810 HCAPLUS Full-text

DOCUMENT NUMBER: 109:193810

TITLE: Organic-electrolyte batteries with improved  
separators

INVENTOR(S): Nishino, Shuichi; Izumikawa, Toshihiko

PATENT ASSIGNEE(S): Matsushita Electric Industrial Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	---	-----	-----	
JP 63155553	A	19880628	JP 1986-302009	198612 18
				198612 18

PRIORITY APPLN. INFO.: JP 1986-302009

AB Separators for organic-**electrolyte** light-metal batteries are prepared by winding a ribbon of a separator material into a tube, fixing the seams, bending 1 end of the tube inwardly to form a cup shape, and inserting a circular piece of nonwoven cloth into the tube to seal the cup-shaped bottom. The ribbon can be nonwoven cloth of synthetic **resin** fibers or **porous** synthetic **resin film**. Thus, separators prepared from nonwoven **polypropylene** cloth tubes with nonwoven **polypropylene** cloth sealing the bottom were used in Li-MnO<sub>2</sub> batteries with 1M LiClO<sub>4</sub>/propylene carbonate-MeOC<sub>2</sub>H<sub>4</sub>OMe **electrolyte**. After 120 heating-cooling cycles between -10° and +60°, none of these batteries showed **electrolyte** leak whereas 10 out of 100 control batteries did. Batteries of the invention had an average internal resistance of 7.5  $\Omega$  with a standard deviation of 1.0  $\Omega$ , vs. 12.6 and 8.6  $\Omega$  for control batteries.

IC ICM H01M002-18  
ICS H01M006-16  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 38  
IT Batteries, **primary**  
(separators, nonwoven polypropylene cloth, manufacture of, for leak  
prevention and low internal resistance)

L26 ANSWER 12 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1988:513480 HCAPLUS Full-text

DOCUMENT NUMBER: 109:113480

TITLE: Lithium battery with porous polymer separator

INVENTOR(S): Okamoto, Osamu; Sasama, Hiroshi; Niso, Kyoshi;  
Iwamaru, Futayasu

PATENT ASSIGNEE(S): Hitachi Maxell, Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 5 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63126159	A	19880530	JP 1986-272402	198611 15
PRIORITY APPLN. INFO.:				JP 1986-272402 198611 15

AB A Li battery has a Li-alloyable metal sheet on the separator side of a Li block with the separator made of **microporous resin film** having pore size  $<0.3 \mu\text{m}$ . The metal is Al, Sn, Zn, Pb, Bi, Si, Sb, and/or Mg. Thus, 2  $25\text{-}\mu\text{m}$  **polypropylene** films having oval pores of  $0.1\text{-}$  and  $0.04\text{-}\mu\text{m}$  diams. were stacked with long diameter of pores on 1 film perpendicular to that on the other film to form a separator for use in a battery having a Li anode block with an Al sheet on the separator side, a CuO-FeS<sub>2</sub> cathode, and a 1M LiClO<sub>4</sub>/2:1 (volume) propylene carbonate-MeOC<sub>2</sub>H<sub>4</sub>OMe **electrolyte**. Batteries of the invention had an internal resistance of 13-20  $\Omega$ , a discharge duration of 560-585 h when discharged through a 2-k $\Omega$  load, and none showed short circuiting after a vibration test (JIS C 5025), vs. the resp. values of 12-18  $\Omega$ , 280-585 h, and 25 short circuiting out of 1000 batteries using a separator of 1  $30\text{-}\mu\text{m}$  **polypropylene** film having  $0.5 + 5.0\text{-}\mu\text{m}$  pores.

IC ICM H01M004-06  
 ICS H01M002-16; H01M004-40  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 38  
 IT Batteries, **primary**  
 (separators, polypropylene, lithium)

L26 ANSWER 13 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 1987:199280 HCAPLUS Full-text  
 DOCUMENT NUMBER: 106:199280  
 TITLE: Nonaqueous batteries  
 INVENTOR(S): Kimura, Yukio  
 PATENT ASSIGNEE(S): Sanyo Electric Co., Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 62055871	A	19870311	JP 1985-195060	198509 04
PRIORITY APPLN. INFO.:				JP 1985-195060 198509 04

AB Title batteries have light-metal (Li or Na or its alloy) anodes, cathodes which expand during discharge, **microporous resin-film** separators, **electrolyte** retaining rings above or below separators, and **electrolyte** layers inside the rings. A button-type battery was prepared by using a Cu oxide-10 graphite-5% fluoro-resin cathode, a Li anode, a microporous **polypropylene** separator, and a **polypropylene** ring filled with **electrolyte** of 1M LiClO<sub>4</sub> in propylene carbonate-MeOC<sub>2</sub>H<sub>4</sub>OMe. This structure prevented the increase of internal resistance due to the removal of **electrolyte** from the separator which was observed in control batteries without the ring.

IC ICM H01M006-16

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 72

IT Batteries, **primary**  
(lithium-copper oxide, nonaq., with **polypropylene** **electrolyte**-retaining rings)

IT 9003-07-0, **Polypropylene**

RL: USES (Uses)

(**electrolyte**-retaining rings, in nonaq. lithium-copper oxide batteries)

L26 ANSWER 14 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1986:504683 HCAPLUS Full-text

DOCUMENT NUMBER: 105:104683

TITLE: Nonaqueous battery

INVENTOR(S): Ubukawa, Satoshi; So, Shinji; Amezutsumi, Toru

PATENT ASSIGNEE(S): Sanyo Electric Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	---	-----	-----	
JP 61077270	A	19860419	JP 1984-197508	198409 20

PRIORITY APPLN. INFO.: JP 1984-197508

198409  
20

AB A nonaq. battery has an anode of Li, Na or their alloys in an anode case, a cathode that expands during discharge in a cathode case, a **porous resin-film** separator, and a conductive ring between the cathode and the cathode case defining a space for **electrolyte** storage. Thus, a mixture of FeS<sub>2</sub> 85, graphite 10, and a fluoro-resin powder 5% was pressed at 2 tons/cm<sup>2</sup> to form a pellet (diameter 11, thickness 1.8 mm), and heated to form a cathode. A stainless steel ring was placed between the cathode and a cathode case, and a 1M LiBF<sub>4</sub>/propylene carbonate-MeOC<sub>2</sub>H<sub>4</sub>OMe **electrolyte** was filled into the storage space. A battery using this cathode structure, a Li anode, and a 0.025-mm-thick porous **polypropylene** separator showed a flat, slowly decreasing discharge voltage for .apprx.600 h when discharged through a 5.6-k $\Omega$  load at 20°, vs. a fast voltage drop after .apprx.200-h discharge for a battery without the ring and **electrolyte** between the cathode and the cathode case.

IC ICM H01M006-16

CC 72-3 (Electrochemistry)

Section cross-reference(s): 52

IT Batteries, **primary**

(button-type, iron sulfide-lithium, with stable discharge voltage)

L26 ANSWER 15 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1967:11278 HCAPLUS Full-text

DOCUMENT NUMBER: 66:11278

TITLE: Effect of reaction of water and electrolytes during their counterdiffusion in polymers

AUTHOR(S): Lobanov, Yu. E.; Shterenzon, A. L.

SOURCE: Lakokrasochnye Materialy i Ikh Primenenie (1966), (5), 42-4

CODEN: LAMAAD; ISSN: 0130-9013

DOCUMENT TYPE: Journal

LANGUAGE: Russian

AB The method usually applied for measuring **electrolyte permeability** through **polymer films** does not consider the influence of counterdiffusion of water. This effect was investigated with high-pressure **polyethylene**, poly(tetrafluoroethylene), and tetrafluoroethylene-vinylidene fluoride copolymer films. The film was clamped between two glass half-cells; one of them was equipped with Pt electrodes and filled with water, the other was filled with HCl or HNO<sub>3</sub> solution. The half-cells were not filled simultaneously, but one after another with a delay of 1-3 hrs. In the 1st stage, the **electrolyte** solution or water penetrated into the dry film, the diffusion being unaffected by counterdiffusion of the other component. In this arrangement, the curves representing the amount of acid passed show a discontinuity corresponding to the effect of incipient counterdiffusion. The rate of acid diffusion decreases from  $4.5 + 10^{-4}$  g./hr. to  $0.5 + 10^{-4}$  for 3.1 cm.<sup>2</sup> poly(tetrafluoroethylene) due to interaction of counter-diffusion of water. Similar results were obtained for other films. When the fronts of penetrating liquids meet, mol. associated structures are formed from mols. of water and **electrolyte**. Thus, light-scattering centers appear in the film.

CC 35 (Synthetic High Polymers)

=>